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1. A composition for administration of a beneficial agent to an organism, comprising:

a solvent mixture, comprising

a hydrophobic solvent; and

a hydrophilic solvent;

a bioerodible polymer; and

a beneficial agent,

wherein the polymer and the beneficial agent are dissolved.

- 2. The composition of claim 1, wherein at least 55 wt% of the solvent mixture is the hydrophobic solvent.
- 3. The composition of claim 1, wherein at least 90 wt% of the solvent mixture is the hydrophobic solvent.
- 4. The composition of claim 1, wherein the hydrophobic solvent has a solubility in water of less than 0.1 wt%.
- 5. The composition of claim 1, wherein the beneficial agent has a concentration from 0.1 mg/ml to 500 mg/ml.
- 6. The composition of claim 1, wherein the beneficial agent has a concentration from 10 mg/ml to 100 mg/ml.
- 7. The composition of claim 1, wherein the composition can be injected through a 25-gauge needle.
- 8. The composition of claim 1, wherein the viscosity of the composition is less than 2000 centipoise.
- 9. The composition of claim 1, wherein less than 25% of the beneficial agent is released in 24 hours following administration *in vivo*.

- 10. The composition of claim 1, wherein the hydrophobic solvent is benzyl benzoate, the hydrophilic solvent is benzyl alcohol, the bioerodible polymer is polylactide, and the beneficial agent is a peptide or protein.
- 11. The composition of claim 1, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a suspension.
- 12. The composition of claim 1, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a solution.
- 13. A composition for administration of a beneficial agent, comprising:

a solvent mixture, comprising
a hydrophobic solvent; and
a hydrophilic solvent;

a bioerodible polymer; and

a beneficial agent,

wherein the viscosity of the composition is less than 2000 centipoise.

- 14. The composition of claim 13 wherein at least 55 wt% of the solvent mixture is the hydrophobic solvent.
- 15. The composition of claim 13, wherein at least 90 wt% of the solvent mixture is the hydrophobic solvent.
- 16. The composition of claim 13, wherein the hydrophobic solvent has a solubility in water of less than 0.1 wt%.
- 17. The composition of claim 13, wherein the composition can be injected through a 28-gauge needle.

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- 18. The composition of claim 13, wherein the composition can be injected through a 30-gauge needle.
- 19. The composition of claim 13, wherein the viscosity of the composition is less than 500 centipoise.
- 20. The composition of claim 13, wherein the hydrophobic solvent is benzyl benzoate, the hydrophilic solvent is benzyl alcohol, the bioerodible polymer is polylactide, and the beneficial agent is a peptide or protein.
- 21. The composition of claim 13, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form a solution.
- 22. The composition of claim 13, wherein the hydrophilic solvent, the hydrophobic solvent, the bioerodible polymer, and the beneficial agent form s suspension.
- 23. A composition for administration of a beneficial agent to an organism, comprising:
- a solvent mixture, the solvent mixture comprising a hydrophobic solvent and a hydrophilic solvent;
 - a bioerodible polymer dissolved in the solvent mixture; and a beneficial agent dissolved in the solvent mixture,
- wherein the viscosity of the composition is less than 2000 centipoise, at least 90 wt% of the solvent mixture is the hydrophobic solvent, the hydrophobic solvent has a solubility in water of less than 0.1 wt%, and less than 25% of the beneficial agent is released in 24 hours following administration *in vivo*.
- 24. The composition of claim 23, wherein the hydrophobic solvent is benzyl benzoate, the hydrophilic solvent is benzyl alcohol, the bioerodible polymer is polylactide, and the beneficial agent is a peptide or protein.

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- 25. A method of administering a beneficial agent, comprising: injecting the composition of claim 1 into an organism through a needle.
- 26. The method of claim 25, wherein the needle is a 25-gauge needle.
- 27. The method of claim 25, wherein the needle is a 28-gauge needle.
- 28. The method of claim 25, wherein the needle is a 30-gauge needle.
- 29. A method of administering a beneficial agent, comprising: injecting the composition of claim 13 into an organism through a needle.
- 30. The method of claim 29, wherein the needle is a 25-gauge needle.
- 31. The method of claim 29, wherein the needle is a 28-gauge needle.
- 32. The method of claim 29, wherein the needle is a 30-gauge needle.
- 33. A method of administering a beneficial agent, comprising: injecting the composition of claim 23 into an organism through a needle.
 - 34. A kit, comprising:

a container;

a hydrophobic solvent;

a hydrophilic solvent;

a bioerodible polymer; and

a beneficial agent,

wherein the amount of said hydrophobic solvent and said hydrophilic solvent is sufficient together to dissolve all of said polymer.

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- 35. The kit of claim 34, comprising a unit dosage of the beneficial agent.
- 36. The kit of claim 34, wherein the hydrophobic solvent, the hydrophilic solvent, and the bioerodible polymer are sterile.
 - 37. The kit of claim 34, further comprising at least one syringe.
 - 38. The kit of claim 34, wherein the container comprises a septum.
 - 39. The kit of claim 37, further comprising at least one needle.
- 40. The kit of claim 39, wherein the beneficial agent, the hydrophobic solvent, the hydrophilic solvent, and the bioerodible polymer, are in said at least one syringe.